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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/506,725	09/04/2004	Daniel W Chan	57203(71699)	7047
21874 7590 03/05/2007 EDWARDS & ANGELL, LLP P.O. BOX 55874 BOSTON, MA 02205		EXAMINER		
			JOYCE, CATHERINE	
			ART UNIT	PAPER NUMBER
			1642	
		,		
SHORTENED STATUTOR	RY PERIOD OF RESPONSE	MAIL DATE	DELIVERY MODE	
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Please find below and/or attached an Office communication concerning this application or proceeding.

If NO period for reply is specified above, the maximum statutory period will apply and will expire 6 MONTHS from the mailing date of this communication.

	Application No.	Applicant(s)			
	10/506,725	CHAN ET AL.			
Office Action Summary	Examiner	Art Unit			
•	Catherine M. Joyce	1642			
The MAILING DATE of this communication app Period for Reply	ears on the cover sheet with the c	orrespondence address			
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DA - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period w - Failure to reply within the set or extended period for reply will, by statute, Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tin vill apply and will expire SIX (6) MONTHS from cause the application to become ABANDONE	N. nely filed the mailing date of this communication. D (35 U.S.C. § 133).			
Status					
1) ☐ Responsive to communication(s) filed on <u>04 Secondary</u> 2a) ☐ This action is FINAL . 2b) ☐ This 3) ☐ Since this application is in condition for alloware closed in accordance with the practice under Expression is the practice of th	action is non-final. nce except for formal matters, pro				
Disposition of Claims					
4) Claim(s) 1-57 is/are pending in the application. 4a) Of the above claim(s) is/are withdraw 5) Claim(s) is/are allowed. 6) Claim(s) is/are rejected. 7) Claim(s) is/are objected to. 8) Claim(s) 1-57 are subject to restriction and/or experience. Application Papers 9) The specification is objected to by the Examine 10) The drawing(s) filed on is/are: a) according and applicant may not request that any objection to the Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the Examine 10.	vn from consideration. election requirement. er. epted or b) □ objected to by the drawing(s) be held in abeyance. Se ion is required if the drawing(s) is ob	e 37 CFR 1.85(a). jected to. See 37 CFR 1.121(d).			
Priority under 35 U.S.C. § 119		.			
12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received.					
Attachment(s) 1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date	4) Interview Summary Paper No(s)/Mail D 5) Notice of Informal I 6) Other:	Pate			

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DETAILED ACTION

. 1. Claims 1-57 are pending.

Election/Restrictions

2. Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

- 1. Claims 1-25, 26, 27-36, as drawn to a method of qualifying breast cancer in a subject comprising (a) measuring at least one biomarker in sample from a subject, and (b) correlating the measurement with breast cancer status, wherein the biomarker is Marker I (BCI) and claims 37-41 as dawn to a kit comprising (a) a capture reagent that binds a biomarker is Marker I (BCI) and (b) a container comprising at least one of the biomarkers.
- Claims 1-25, 26, 27-36, as drawn to a method of qualifying breast cancer in a subject comprising (a) measuring at least one biomarker in sample from a subject, and (b) correlating the measurement with breast cancer status, wherein the biomarker is Marker II (BC2), Marker III (BC3), Marker IV, Marker V, Marker VI, Marker VII, Marker VIII, Marker XI, Marker XI, Marker XIII, Marker XIV.
- Claims 37-41, as drawn to a kit comprising (a) a capture reagent that binds a biomarker is selected from the group consisting of Marker II (BC2), Marker III (BC3), Marker IV, Marker V, Marker VI, Marker VII, Marker VIII,

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Marker IX, Marker X, Marker XI, Marker XII, Marker XIII, Marker XIV and (b) a container comprising at least one of the biomarkers.

- 4. Claims 42-46, as drawn to a kit comprising (a) a first capture reagent that binds at least one biomarker selected from the group consisting of Marker I (BCI), Marker II (BC2), Marker III (BC3), Marker IV, Marker V, Marker VI, Marker VII, Marker VIII, Marker IX, Marker X, Marker XI, Marker XIII, Marker XIV, and (b) a second capture reagent that binds at least one of the biomarkers that is not bound by the first capture reagent.
- 5. Claims 47-53, as drawn to a kit comprising (a) a first capture reagent that binds at least one biomarker selected from the group consisting of Marker I (BCI), Marker II (BC2), Marker III (BC3), Marker IV, Marker V, Marker VI, Marker VII, Marker VIII, Marker IX, Marker X, Marker XI, Marker XIII, Marker XIV, and (b) instructions for using the capture reagent to detect the biomarker.
- 6. Claims 54-56, as drawn to a kit comprising (a) a first capture reagent that binds at least one biomarker that binds to at least two biomarkers selected from the group consisting of selected from the group consisting of Marker I (BCI), Marker II (BC2), Marker III (BC3), Marker IV, Marker V, Marker VI, Marker VII, Marker VIII, Marker IX, Marker X, Marker XI, Marker XII, Marker XIII, Marker XIV, and (b) instructions for using the capture reagent to detect the biomarker.
- 7. Claims 57, as drawn to a kit comprising (a) a plurality of capture reagents each of which has bound to it a different biomarker selected from the group consisting of Marker I (BCI), Marker II (BC2), Marker III (BC3), Marker IV, Marker V, Marker VI, Marker VII, Marker VIII, Marker VIII, Marker XI, Marker XI, Marker XIII, Marker XIV.

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3. The inventions are distinct, each from the other, because of the following reasons:

The inventions listed as Groups 1-7 do not relate to a single inventive concept because they lack the same or corresponding special technical features for the following reasons:

An international stage application shall relate to one invention only or to a group of inventions so linked as to form a single general inventive concept. When claims to different categories are present in the application, the claims will be considered to have unity of invention if the claims are drawn only to one of the following combinations of categories: (1) A product and a process specially adapted for the manufacture of said product; or (2) A product and a process of use of said product; or (3) A product, a process specially adapted for the manufacture of the said product, and a use of the said product; or (4) A product, a process specially adapted for the manufacture of the said product, and an apparatus or means specifically designed for carrying out the said process; or (5) A product, a process specially adapted for the manufacture of the said product, and an apparatus or means specifically designed for carrying out the said process. If multiple products, processes of manufacture or uses are claimed, the first invention of the category first mentioned in the claims of the application will be considered as the main invention in the claims, see PCT article 17(3) (a) and 1.476 (c), 37 C.F.R.

1.475(b) and (d). Group I will be the main invention. After that, all other products and methods will be broken out as separate groups (see 37 CFR 1.475(d).)

4. Further, the following elections of species are required.

If Groups 1 or 2 are elected, election of a specific managing treatment step from the following list is required: ordering more rests, performing surgery, and taking no further action.

If Groups 1 or 2 are elected, election of a specific breast cancer status from the following list is required: the subject's risk of cancer, the presence or absence of disease, the stage of disease, and the effectiveness of treatment.

If Groups 1, 2, 3 or 6 are elected, election of a specific known breast cancer marker from the following list is required: CA 15.3 or CA 27.29.

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If Groups 1 or 2 are elected, election of a specific substrate from from the following list is required: a SELDI probe comprising an IMAC Ni surface; a SELDI probe comprising biospecific affinity reagents; a microtiter plate comprising biospecific affinity reagents.

If Groups 1 or 2 are elected, election of a specific measuring type in the following list is required: detecting the presence or absence of the biomarkers; quantifying the amount of marker(s); qualifying the type of biomarker.

If Groups 1 or 2 are elected, election of a specific biochip array type from the following list is required: a nucleic acid array; a protein chip array.

If Groups 1 or 2 are elected, election of specific method of protein detection from the following list is required: SELDI or immunoassay.

If Group 2 is elected, election of a specific marker from the following list is required: Marker II (BC2), Marker III (BC3), Marker IV, Marker V, Marker VI, Marker VII, Marker XI, Marker XII, Marker XIV.

If Groups 1, 3 or 6 is elected, election of second biomarker (different from the first biomarker) from the following list is required: Marker I (BCI), Marker II (BC2), Marker III (BC3), Marker IV, Marker V, Marker VI, Marker VII, Marker VIII, Marker IX, Marker X, Marker XII, Marker XIII, Marker XIV

If Groups 3 is elected, election of a biomarker from the following list is required: Marker II (BC2), Marker III (BC3), Marker IV, Marker V, Marker VI, Marker VIII, Marker XI, Marker XII, Marker XIII, Marker XIV.

If Groups 4-7 are elected, election of a biomarker from the following list is required: Marker I (BCI), Marker II (BC2), Marker III (BC3), Marker IV, Marker V, Marker VI, Marker VII, Marker VII, Marker IX, Marker X, Marker XI, Marker XIII, Marker XIV.

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If Groups 4-5 are elected, election of a specific capture reagent from the following list is required: an antibody; an immobilized metal chelate.

If Groups 4-5 are elected, election of a specific capture reagent from the following list is required: an antibody; an immobilized metal chelate.

- 5. Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP 809.02(a).
- 6. Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C.103 of the other invention.
- This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103, the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 C.F.R. 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of potential 35 U.S.C. 102(f) or (g) prior art under 35 U.S.C. 103.
- 8. Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim

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remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

9. Applicant is advised that the response to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed.

10. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Catherine M. Joyce whose telephone number is 571-272-3321. The examiner can normally be reached on Monday thru Friday, 10:15 - 6:45.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Shanon Foley, can be reached on 571-272-0898. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8700.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

- (1-14A11 1010) 77, 1014.D

Catherine M. Joyce Examiner

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